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| **UK Renal Registry data request – expression of interest** | |
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| Thank you for your interest in using pseudonymised patient level data held by the Renal Association’s (RA’s) UK Renal Registry (UKRR). To ensure as smooth an application process as possible, we need to understand the data you wish to obtain and how you intend using it. To help us with this, we ask that you complete this short expression of interest (EOI) and email it, along with any supporting documents, to  [ukrr-research@renalregistry.nhs.uk](mailto:ukrr-research@renalregistry.nhs.uk)  Currently, we only release patient level data to academic and clinical organisations – we do not release patient level data to commercial organisations.  If you would like to work with the UKRR on a grant application, please email [ukrr-research@renalregistry.nhs.uk](mailto:ukrr-research@renalregistry.nhs.uk). Once you have secured funding we will ask you to apply through the data request process – until then, you do not need to complete this form.  You can submit your EOI at any time. It will be assessed by a review panel comprising members with diverse skill-sets. It is very important, therefore, that all questions are answered in **clear and plain** English.  If your EOI is approved by the review panel, you will be invited to complete a full application form and data protection impact assessment (DPIA). Your full application and DPIA will be assessed by clinical and methodological experts who meet on a quarterly basis as the RA Data Release Group – please ensure you submit your application and DPIA at least two weeks prior to the meeting in which you want them to be reviewed – for meeting dates see [renal.org/audit-research/how-access-data/ukrr-data/apply-access-ukrr-data](https://renal.org/audit-research/how-access-data/ukrr-data/apply-access-ukrr-data).    If your application and DPIA are approved, you will be asked to complete a data sharing agreement (DSA) with the RA prior to data release. | |
| **RA use only** | |
| Application number |  |
| Date received |  |
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| **Applicant, please complete all the following fields** | |
| Project title |  |
| Main applicant |  |
| EOI author, if different to main applicant |  |
| Email address |  |
| Telephone no. |  |
| Institution/organisation |  |
| Co-applicants, including their email addresses |  |
| Will you be collaborating with the commercial sector? If so, provide details |  |
| Is the project funded? If so, provide funder details |  |
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| Project summary, addressing each of the bullet points (max 400 words in plain English) – this will be shared with the Renal Association Patient Council, so please write your responses below each of the bullet points | |
| * What is already known about this topic and why is it important? * How will you carry out your study? * How will you decide which patients are included in your study? * How many patients do you anticipate including? * For how long will you follow up these patients? * What value will UKRR data add to the project? * What new information will your study generate and how will this benefit patients? | |
| Project objectives (max 100 words) | |
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| Key deliverables, including outputs (max 100 words) | |
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| What is the age range of your cohort? | Paediatric, i.e. <18 years |  |
| Adult, i.e. >18 years |  |
| Paediatric and adult |  |

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| Is your project research or non-research?  For guidance, complete the four questions of the [Health Research Authority decision tool](http://www.hra-decisiontools.org.uk/research/) – leave the IRAS project ID field blank | | Research | |  |
| Non-research | |  |
|  | | | | |
| What permissions do you have in place to conduct your research? Will you need to apply for ethics permission from your institution? | |  | | |
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| Does your team have a statistician to conduct the required analyses? It might also be useful to have someone on your team with epidemiological expertise to help interpret the data | | Yes, we have a statistician | |  |
| Yes, we have a statistician and epidemiologist | |  |
| No, we do not have a statistician | |  |
| No, we would like to buy a UKRR statistician’s time | |  |
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| What UKRR-held data items would you like to obtain? Please check data completeness in the [UKRR data portal](https://renal.org/audit-research/data-portal)  If you are interested in accessing data items held in RaDaR, PatientView and/or in the shared NHSBT-UKRR dataset, please contact the research team directly, because different permissions and processes apply. Also, please note that the UKRR is unable to release any comorbidity data from HES – you must apply yourself | | | | |
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| Are you planning to link UKRR data to other data? (If yes, please state the dataset(s) you intend to link to) | |  | | |
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| Please return your completed expression of interest to [ukrr-research@renalregistry.nhs.uk](mailto:ukrr-research@renalregistry.nhs.uk) | | | | |